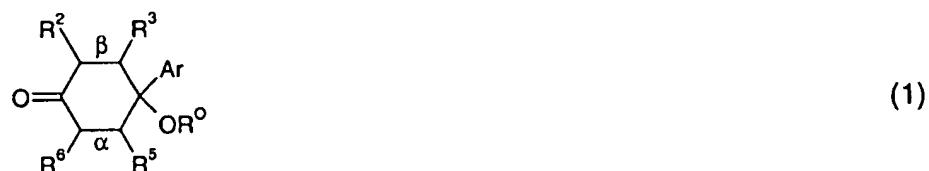


AMENDMENTS TO THE CLAIMS

Amend the claimed as follows:

Claims 1-89. (Cancelled)

90. (new) A compound having the following formula:



wherein:

Ar is a 1-(sulfonyl)-1H-indol-2-yl group;

the group -OR^O is independently:

- (a) -OH;
- (b) an ether group; or:
- (c) an acyloxy group;

the bond marked α is independently:

- (a) a single bond; or:
- (b) a double bond;

the bond marked β is independently:

- (a) a single bond; or:
- (b) a double bond;

each of R², R³, R⁵, and R⁶, is independently a ring substituent and is:

- (a) H;
- (b) a monovalent monodentate substituent; or:

(c) a ring substituent which, together with an adjacent ring substituent, and together with the ring atoms to which these ring substituents are attached, form a fused ring;

and pharmaceutically acceptable salts, esters, amides, solvates, hydrates, and protected forms thereof.

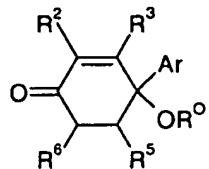
91. (new) A compound according to claim 90, wherein α is independently a double bond and β is independently a double bond, and the compound has the following formula:



92. (new) A compound according to claim 90, wherein α is independently a single bond and β is independently a single bond and the compound has the following formula:



93. (new) A compound according to claim 90, wherein α is independently a single bond and β is independently a double bond, and the compound has the following formula:



94. (new) A compound according to claim 90, wherein said monovalent monodentate substituent is selected from:

hydroxy (-OH);

halo;

cyano (-CN);

carboxy (-COOH);

azido;

ester;

amino, including:

C₁₋₇alkyl-amino;

amino-C₁₋₇alkyl-amino;

C₁₋₇alkyl, including:

halo-C₁₋₇alkyl;

amino-C₁₋₇alkyl;

carboxy-C₁₋₇alkyl;

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hydroxy-C₁₋₇alkyl;
C₅₋₂₀aryl-C₁₋₇alkyl;
ether, including:
 C₁₋₇alkoxy;
 halo-C₁₋₇alkoxy;
 amino-C₁₋₇alkoxy;
 carboxy-C₁₋₇alkoxy;
 hydroxy-C₁₋₇alkoxy;
 C₅₋₂₀aryl-C₁₋₇alkoxy;
acyl, including:
 C₁₋₇alkyl-acyl;
 halo-C₁₋₇alkyl-acyl;
 amino-C₁₋₇alkyl-acyl;
 carboxy-C₁₋₇alkyl-acyl;
 hydroxy-C₁₋₇alkyl-acyl;
 C₅₋₂₀aryl-C₁₋₇alkyl-acyl;
C₅₋₂₀aryl-acyl;
C₅₋₂₀aryl;
thiol (-SH); and,
thioether.

95. (new) A compound according to claim 90, wherein said monovalent monodentate substituent is selected from:

- OH;
- F, -Cl, -Br, -I;
- CN;
- COOH;
- N₃;
- COOMe, -COOEt, -COOtBu, -COOPh, -COOCH₂Ph;

- NH₂, -NHMe, -NHEt, -NMe₂, -NEt₂;
- piperidino, morpholino, piperazino, N-methyl-piperazino;
- NH(CH₂)_w-NH₂, -NH(CH₂)_w-NHMe, -NH(CH₂)_w-NMe₂, -NH(CH₂)_w-NEt₂;

- Me, -Et, -nPr, -iPr, -nBu, -iBu, -sBu, -tBu;
- CH₂F, -CH₂Cl, -CF₃, -CCl₃, -CF₂CF₃, -CH₂CF₃, -C(CF₃)₃;
- (CH₂)_w-NH₂, -(CH₂)_w-NHMe, -(CH₂)_w-NMe₂, -(CH₂)_w-NEt₂;
- (CH₂)_w-COOH;
- (CH₂)_w-OH;
- CH₂Ph;

- OMe, -OEt, -OnPr, -OiPr, -OnBu, -OiBu, -OsBu, -OtBu;

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-OCH₂F, -OCH₂Cl, -OCF₃, -OCCl₃, -OCF₂CF₃, -OCH₂CF₃, -OC(CF₃)₃;
-O(CH₂)_w-NH₂, -O(CH₂)_w-NHMe, -O(CH₂)_w-NMe₂, -O(CH₂)_w-NEt₂;
-O(CH₂)_w-COOH;
-O(CH₂)_w-OH;
-OCH₂Ph;

-C(=O)Me, -C(=O)Et, -C(=O)-nPr, -C(=O)-iPr, -C(=O)-nBu, -C(=O)-iBu,
-C(=O)-sBu, -C(=O)-tBu;
-C(=O)CH₂F, -C(=O)CH₂Cl, -C(=O)CF₃, -C(=O)CCl₃, -C(=O)CF₂CF₃,
-C(=O)CH₂CF₃, -C(=O)C(CF₃)₃;
-C(=O)(CH₂)_w-NH₂, -C(=O)(CH₂)_w-NHMe, -C(=O)(CH₂)_w-NMe₂,
-C(=O)(CH₂)_w-NEt₂;
-C(=O)(CH₂)_w-COOH;
-C(=O)(CH₂)_w-OH;
-C(=O)CH₂Ph;

-Ph;

-SH;
-SMe, -SEt, -SnPr, -S-iPr, -S-nBu, -S-iBu, -S-sBu, -S-tBu,
-S-CH₂-Ph, -S-Ph;

a thioether group derived from cysteine, homocysteine, glutathione, or a peptide comprising the sequence -Cys-(X)_y-Cys-, where X is an amino acid, and y is an integer from 1 to 6;

wherein w is an integer from 1 to 7.

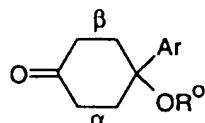
96. (new) A compound according to claim 90, wherein each of R², R³, R⁵, and R⁶, is independently a ring substituent and is:

- (a) H; or:
- (b) a monovalent monodentate substituent.

97. (new) A compound according to claim 91, wherein each of R², R³, R⁵, and R⁶, is independently a ring substituent and is:

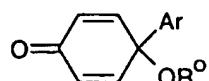
- (a) H; or:
- (b) a monovalent monodentate substituent.

98. (new) A compound according to claim 90, wherein R², R³, R⁵ and R⁶ are -H:



(9)

99. (new) A compound according to claim 90, wherein R², R³, R⁵ and R⁶ are -H; α is a double bond; and β is a double bond:

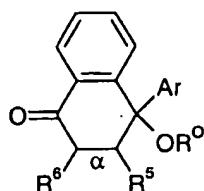


(11)

100. (new) A compound according to claim 90, wherein

- (a) R² and R³, together with the ring atoms to which they are attached, form a fused ring; or
- (b) R⁵ and R⁶, together with the ring atoms to which they are attached, form a fused ring; or
- (c) or both (a) and (b).

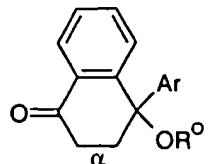
101. (new) A compound according to claim 99, wherein R² and R³ form a fused benzene ring; and β is a double bond:



(14)

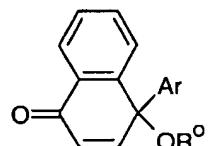
102. (new) A compound according to claim 101, wherein R⁵ and R⁶ do not also form a fused ring.

103. (new) A compound according to claim 99, wherein R² and R³ form a fused benzene ring; β is a double bond; and R⁵ and R⁶ are -H:



(17)

104. (new) A compound according to claim 99, wherein R² and R³ form a fused benzene ring; β is a double bond; R⁵ and R⁶ are -H; and α is a double bond:



(18)

105. (new) A compound according to claim 90, wherein R⁰ is independently:

(a) -H;

(b) C₁₋₇alkyl, C₃₋₂₀heterocyclyl, or C₅₋₂₀aryl; and is optionally substituted;

or:

(c) C₁₋₇alkyl-acyl, C₃₋₂₀heterocyclyl-acyl, or C₅₋₂₀aryl-acyl; and is optionally substituted.

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106. (new) A compound according to claim 104, wherein R^O is optionally substituted with one more of the following groups:

hydroxy (-OH);

halo;

carboxy (-COOH);

amino; and,

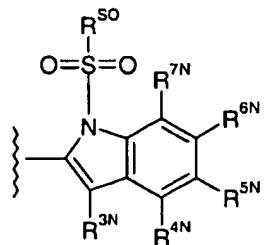
C₅₋₂₀aryl.

107. (new) A compound according to claim 90, wherein R^O is -H.

108. (new) A compound according to claim 91, wherein R^O is -H.

109. (new) A compound according to claim 99, wherein R^O is -H.

110. (new) A compound according to claim 90, wherein Ar is a group of the following formula:



wherein:

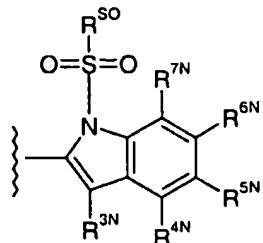
R^{SO} is independently a sulfonyl substituent; and

each of R^{3N} , R^{4N} , R^{5N} , R^{6N} , and R^{7N} is independently an indolyl substituent.

111. (new) A compound according to claim 110, wherein R^{SO} is C_{1-7} alkyl, C_{3-20} heterocyclyl, or C_{5-20} aryl; and is optionally substituted.

112. (new) A compound according to claim 110, wherein R^{SO} is C_{5-20} aryl; and is optionally substituted.

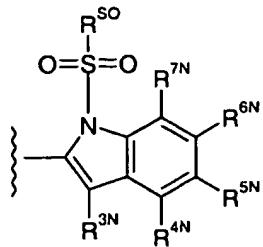
113. (new) A compound according to claim 99, wherein Ar is a group of the following formula:



wherein:

R^{SO} is independently C_{5-20} aryl; and is optionally substituted; and each of R^{3N} , R^{4N} , R^{5N} , R^{6N} , and R^{7N} is independently an indolyl substituent.

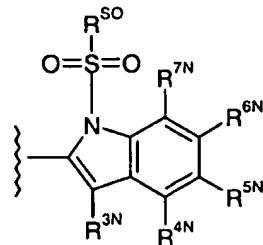
114. (new) A compound according to claim 104, wherein Ar is a group of the following formula:



wherein:

R^{SO} is independently C₅₋₂₀aryl; and is optionally substituted; and each of R^{3N}, R^{4N}, R^{5N}, R^{6N}, and R^{7N} is independently an indolyl substituent.

115. (new) A compound according to claim 109, wherein Ar is a group of the following formula:



wherein:

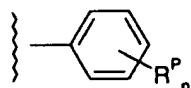
R^{SO} is independently C₅₋₂₀aryl; and is optionally substituted; and each of R^{3N}, R^{4N}, R^{5N}, R^{6N}, and R^{7N} is independently an indolyl substituent.

116. (new) A compound according to claim 110, wherein R^{SO} is phenyl or naphthyl; and is optionally substituted.

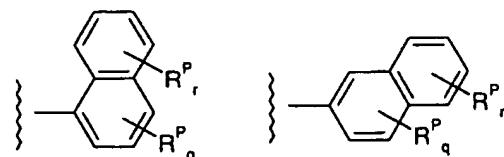
117. (new) A compound according to claim 110, wherein R^{SO} is naphthyl; and is optionally substituted.

118. (new) A compound according to claim 110, wherein R^{SO} is phenyl; and is optionally substituted.

119. (new) A compound according to claim 110, wherein R^{SO} is selected from:



wherein p is an integer from 0 to 5, and each R^P is a phenyl substituent;
and



wherein q is an integer from 0 to 3; r is an integer from 0 to 4; and each R^P is a naphthyl substituent.

120. (new) A compound according to claim 119, wherein each R^P is independently selected from:

hydroxy (-OH);

halo;

cyano (-CN);

carboxy (-COOH);

azido;

ester;

amino, including:

amino-C₁₋₇alkyl-amino;

C₁₋₇alkyl, including:

halo-C₁₋₇alkyl;

amino-C₁₋₇alkyl;

carboxy-C₁₋₇alkyl;

hydroxy-C₁₋₇alkyl;

C₅₋₂₀aryl-C₁₋₇alkyl;

ether, including:

C₁₋₇alkoxy;

halo-C₁₋₇alkoxy;

amino-C₁₋₇alkoxy;

carboxy-C₁₋₇alkoxy;

hydroxy-C₁₋₇alkoxy;

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C₅₋₂₀aryl-C₁₋₇alkoxy;
acyl, including:
C₁₋₇alkyl-acyl;
halo-C₁₋₇alkyl-acyl;
amino-C₁₋₇alkyl-acyl;
carboxy-C₁₋₇alkyl-acyl;
hydroxy-C₁₋₇alkyl-acyl;
C₅₋₂₀aryl-C₁₋₇alkyl-acyl;
C₅₋₂₀aryl-acyl;
C₅₋₂₀aryl.

121. (new) A compound according to claim 119, wherein each R^P is independently selected from:

-OH;
-F, -Cl, -Br, -I;
-CN;
-COOH;
-N₃;
-COOMe, -COOEt, -COOtBu, -COOPh, -COOCH₂Ph;

-NH₂, -NHMe, -NHEt, -NMe₂, -NEt₂;

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piperidino, morpholino, piperazino, N-methyl-piperazino;
-NH(CH₂)_w-NH₂, -NH(CH₂)_w-NHMe, -NH(CH₂)_w-NMe₂, -NH(CH₂)_w-NEt₂;

-Me, -Et, -nPr, -iPr, -nBu, -iBu, -sBu, -tBu;
-CH₂F, -CH₂Cl, -CF₃, -CCl₃, -CF₂CF₃, -CH₂CF₃, -C(CF₃)₃;
-(CH₂)_w-NH₂, -(CH₂)_w-NHMe, -(CH₂)_w-NMe₂, -(CH₂)_w-NEt₂;
-(CH₂)_w-COOH;
-(CH₂)_w-OH;
-CH₂Ph;

-OMe, -OEt, -OnPr, -OiPr, -OnBu, -OiBu, -OsBu, -OtBu;
-OCH₂F, -OCH₂Cl, -OCF₃, -OCCl₃, -OCF₂CF₃, -OCH₂CF₃, -OC(CF₃)₃;
-O(CH₂)_w-NH₂, -O(CH₂)_w-NHMe, -O(CH₂)_w-NMe₂, -O(CH₂)_w-NEt₂;
-O(CH₂)_w-COOH;
-O(CH₂)_w-OH;
-OCH₂Ph;

-C(=O)Me, -C(=O)Et, -C(=O)-nPr, -C(=O)-iPr, -C(=O)-nBu, -C(=O)-iBu,
-C(=O)-sBu, -C(=O)-tBu;
-C(=O)CH₂F, -C(=O)CH₂Cl, -C(=O)CF₃, -C(=O)CCl₃, -C(=O)CF₂CF₃,
-C(=O)CH₂CF₃, -C(=O)C(CF₃)₃;

-C(=O) (CH₂)_w-NH₂, -C(=O) (CH₂)_w-NHMe, -C(=O) (CH₂)_w-NMe₂,
-C(=O)(CH₂)_w-NEt₂;
-C(=O) (CH₂)_w-COOH;
-C(=O) (CH₂)_w-OH;
-C(=O)CH₂Ph;

-Ph;

wherein w is an integer from 1 to 7.

122. (new) A compound according to claim 119, wherein each R^P is independently selected from: -F, -Cl, -Br, -I, -Me, -Et, -OMe, -OEt.

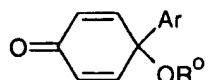
123. (new) A compound according to claim 119, wherein each R^P is independently selected from: -F, -Me, -OMe.

124. (new) A compound according to claim 120, wherein each of R^{3N}, R^{4N}, R^{5N}, R^{6N}, and R^{7N} is independently -H, or as defined for R^P.

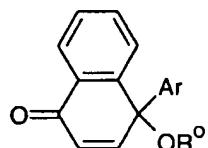
125. (new) A compound according to claim 120, wherein each of R^{3N}, R^{4N}, R^{5N}, R^{6N}, and R^{7N} is independently selected from:
-H, -F, -Cl, -Br, -I, -Me, -Et, -OMe, -OEt.

126. (new) A compound according to claim 120, wherein each of R^{3N}, R^{4N}, R^{6N}, and R^{7N} is -H.

127. (new) A compound selected from compounds having the following formulae and pharmaceutically acceptable salts, esters, amides, solvates, hydrates, and protected forms thereof:



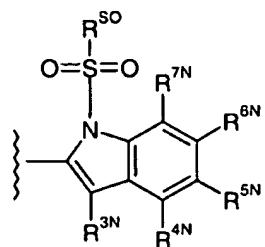
(11)



(18)

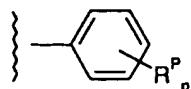
wherein R^o is -H;

wherein Ar is a group of the following formula:

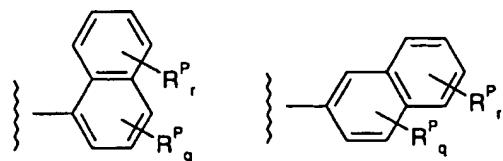


wherein:

R^{SO} is selected from:



wherein p is an integer from 0 to 5, and each R^P is a phenyl substituent; and



wherein q is an integer from 0 to 3; r is an integer from 0 to 4; and each R^P is a naphthyl substituent;

and wherein:

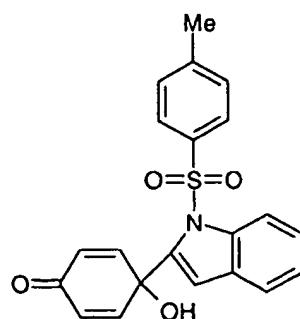
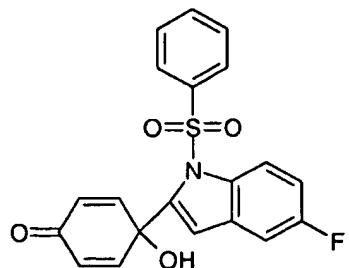
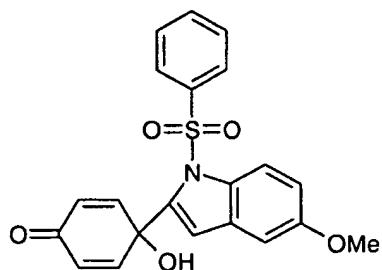
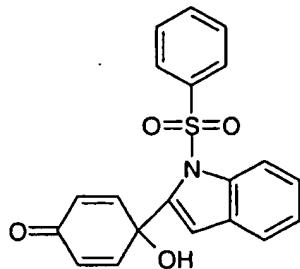
each of R^{3N} , R^{4N} , R^{5N} , R^{6N} , and R^{7N} is independently an indolyl substituent.

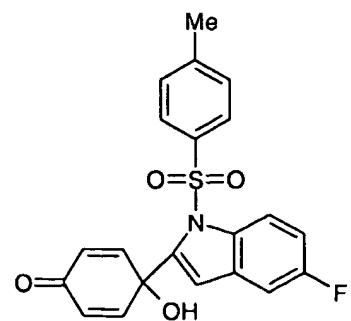
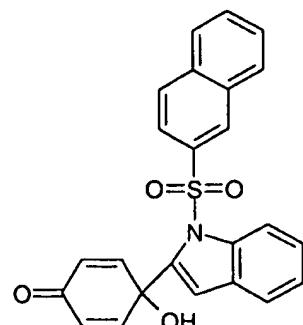
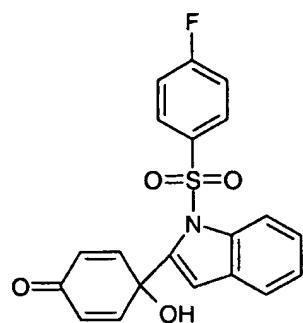
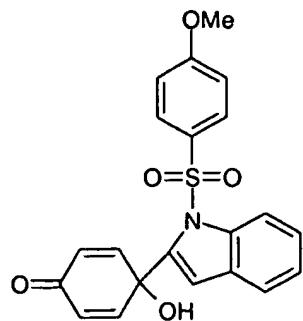
128. (new) A compound according to claim 127, wherein:

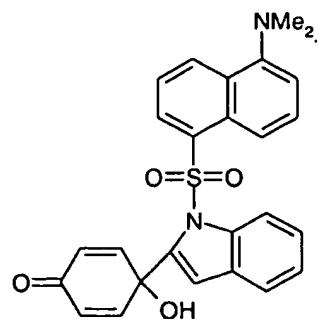
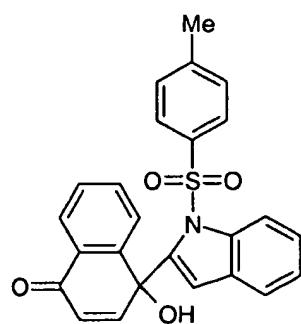
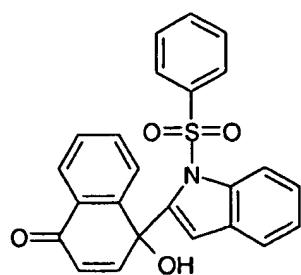
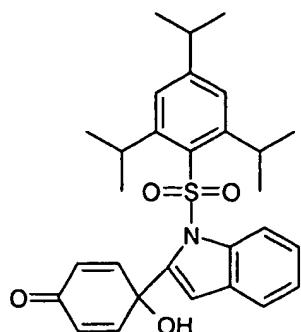
each R^P is independently selected from: -F, -Cl, -Br, -I, -Me, -Et, -OMe, -OEt; and

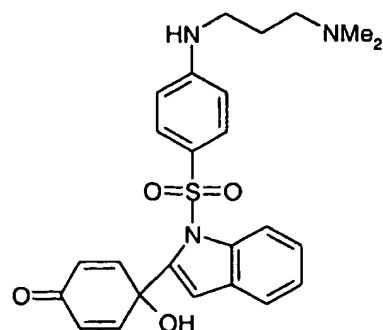
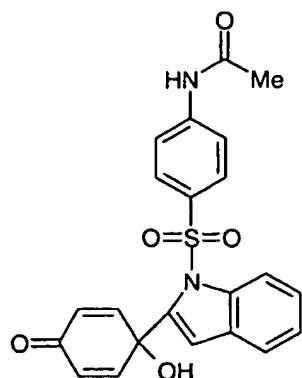
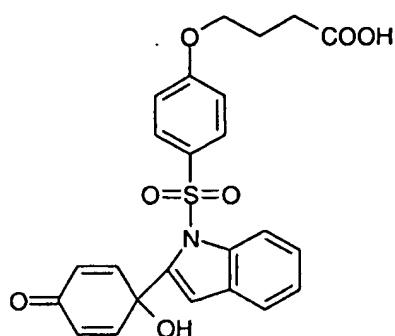
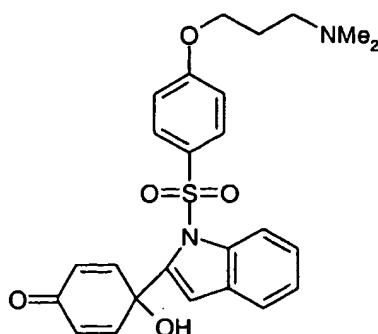
each of R^{3N} , R^{4N} , R^{5N} , R^{6N} , and R^{7N} is independently selected from: -H, -F, -Cl, -Br, -I, -Me, -Et, -OMe, -OEt.

129. (new) A compound selected from the following compounds and pharmaceutically acceptable salts, esters, amides, solvates, hydrates, and protected forms thereof:









130. (new) A composition comprising a compound according to claim 90 and a pharmaceutically acceptable carrier or diluent.

131. (new) A method for the treatment of a proliferative condition comprising administering to a subject suffering from said condition a therapeutically-effective amount of a compound according to claim 90.

132. (new) A method for the treatment of cancer comprising administering to a subject suffering from said cancer a therapeutically-effective amount of a compound according to claim 90.

133. (new) A method for the treatment of colon cancer or renal cancer comprising administering to a subject suffering from said cancer a therapeutically-effective amount of a compound according to claim 90.

134. (new) A method for the treatment of a condition mediated by thioredoxin/thioredoxin reductase comprising administering to a subject suffering from said condition a therapeutically-effective amount of a compound according to claim 90.

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135. (new) A method of inhibiting thioredoxin/thioredoxin reductase in a cell, *in vitro* or *in vivo*, comprising contacting said cell with an effective amount of according to claim 90.

136. (new) A method of regulating cell proliferation, *in vitro* or *in vivo*, comprising contacting a cell with an effective amount of a compound according to claim 90.

137. (new) A method of (a) inhibiting cell proliferation; (b) inhibiting cell cycle progression; (c) promoting apoptosis; or (d) a combination of one or more of these, *in vitro* or *in vivo*, comprising contacting a cell with an effective amount of a compound according to claim 90.